

FORM PTO-1390 REV. 5-93		US DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTORNEYS DOCKET NUMBER P00,0164
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371			U.S. APPLICATION NO. (if known, see 37 CFR 1.5) 09/508635
INTERNATIONAL APPLICATION NO. PCT/EP98/05843	INTERNATIONAL FILING DATE 14 September 1998	PRIORITY DATE CLAIMED 16 September 1997	
TITLE OF INVENTION "ORGAN SPECIFIC NUTRITION"			
APPLICANT(S) FOR DO/EO/US Olivier BALLEVRE, Lionel BOVETTO, Sophie CHARRIER-BROMONT, Jean GRIZARD, and Jean-Claude MAIRE			
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:			
<p>1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.</p> <p>2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.</p> <p>3. <input checked="" type="checkbox"/> This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay.</p> <p>4. <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.</p> <p>5. <input checked="" type="checkbox"/> A copy of International Application as filed (35 U.S.C. 371(c)(2))</p> <p>a. <input checked="" type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau)</p> <p>b. <input type="checkbox"/> has been transmitted by the International Bureau.</p> <p>c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US)</p> <p>6. <input type="checkbox"/> A translation of the International Application into English (35 U.S.C. 371(c)(2)).</p> <p>7. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. §371(c)(3))</p> <p>a. <input checked="" type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau).</p> <p>b. <input type="checkbox"/> have been transmitted by the International Bureau.</p> <p>c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired.</p> <p>d. <input type="checkbox"/> have not been made and will not be made.</p> <p>8. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).</p> <p>9. <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)). (unexecuted)</p> <p>10. <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).</p> <p>Items 11. to 16. below concern other document(s) or information included:</p> <p>11. <input type="checkbox"/> An Information Disclosure Statement under 37 C.F.R. 1.97 and 1.98; (PTO 1449, Prior Art, Search Report).</p> <p>12. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 C.F.R. 3.28 and 3.31 is included. (SEE ATTACHED ENVELOPE)</p> <p>13. <input type="checkbox"/> A FIRST preliminary amendment. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment.</p> <p>14. <input type="checkbox"/> A substitute specification.</p> <p>15. <input type="checkbox"/> A change of power of attorney and/or address letter.</p> <p>16. <input checked="" type="checkbox"/> Other items or information:</p> <p>▪ EXPRESS MAIL #EI 428 568 614 US, dated March 14, 2000.</p>			

U.S. APPLICATION NO. (if known, see 37 C.F.R. 1.5) 09/508635		INTERNATIONAL APPLICATION NO. PCT/EP98/05843		ATTORNEY'S DOCKET NUMBER P00,0164	
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17. <input checked="" type="checkbox"/> The following fees are submitted:				CALCULATIONS	PTO USE ONLY
BASIC NATIONAL FEE (37 C.F.R. 1.492(a)(1)-(5)): Search Report has been prepared by the EPO or JPO \$840.00 International preliminary examination fee paid to USPTO (37 C.F.R. 1.482) \$670.00 No international preliminary examination fee paid to USPTO (37 C.F.R. 1.482) but international search fee paid to USPTO (37 C.F.R. 1.445(a)(2)) \$760.00 Neither international preliminary examination fee (37 C.F.R. 1.482) nor international search fee (37 C.F.R. 1.445(a)(2)) paid to USPTO \$970.00 International preliminary examination fee paid to USPTO (37 C.F.R. 1.482) and all claims satisfied provisions of PCT Article 33(2)-(4) \$ 96.00					
ENTER APPROPRIATE BASIC FEE AMOUNT =					
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 C.F.R. 1.492(e)).				\$	
Claims	Number Filed	Number Extra	Rate		
Total Claims	10 - 20 =		X \$ 18.00	\$	
Independent Claims	1 - 3 =		X \$ 78.00	\$	
Multiple Dependent Claims			\$260.00 +	\$	
TOTAL OF ABOVE CALCULATIONS =				\$ 840.00	
Reduction by 1/2 for filing by small entity, if applicable. Verified Small Entity statement must also be filed. (Note 37 C.F.R. 1.9, 1.27, 1.28)				\$	
SUBTOTAL =				\$	
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).				\$	
TOTAL NATIONAL FEE =				\$	
Fee for recording the enclosed assignment (37 C.F.R. 1.21(h). The assignment must be accompanied by an appropriate cover sheet (37 C.F.R. 3.28, 3.31). \$40.00 per property				\$	
TOTAL FEES ENCLOSED =				\$ 840.00	
				Amount to be refunded	\$
				charged	\$

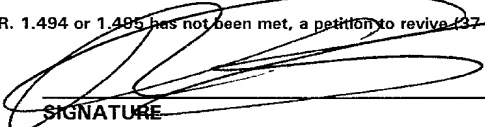
a. ☒ A check in the amount of \$ 840.00 to cover the above fees is enclosed.

b. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 08-2290. A duplicate copy of this sheet is enclosed.

NOTE: Where an appropriate time limit under 37 C.F.R. 1.494 or 1.495 has not been met, a petition to revive (37 C.F.R. 1.137(a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

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 SIGNATURE

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 REGISTRATION NUMBER

Organ Specific Nutrition

Field of the Invention

This invention relates to a method of providing nutrition to a patient for causing a desired effect in a selected organ; for example increasing the protein concentration or rate of protein synthesis in the organ.

Background to the Invention

Nutritional formulas based upon naturally occurring proteins are well known in infant nutrition and clinical nutrition; especially formulas based upon milk and soy proteins. Further, hydrolysates of these proteins are commonly used in infant nutrition and clinical nutrition; particularly in hypoallergenic formulas and formulas for patients suffering from various intestinal absorption problems. Also, hydrolysates are commonly used in clinical nutrition due to their reduced tendency, as compared to proteins, to coagulate. It is also known to use free amino acids in nutritional formulas; either alone or in combination with protein or protein hydrolysates. Usually free amino acids are used for patients suffering from particular diseases or conditions such as inflammatory bowel disease, intractable diarrhoea, short bowel syndrome, and the like.

In all cases, the protein, hydrolysates or free amino acids in the nutritional formulas are intended to provide a source of amino acids to meet the general amino acid needs of the patient. Whether protein, hydrolysate or free amino acid, or a mixture of any of these is used, usually depends upon the condition of the intended consumer. If the intended consumer is a normal, healthy person, whole dietary protein is generally used. However, if the person suffers from a particular disease or condition, or is allergic to whole protein or is at risk of developing an allergy, a hydrolysate or free amino acid mix which the person is better able to tolerate or absorb is generally used.

There is also an interest in using protein hydrolysates in nutrition since it is generally accepted that protein hydrolysates are more rapidly absorbed in the intestine than whole protein or free amino acids (Rerat A.A. ; 1993; Proceedings of the Nutrition Society, 52, 335-344). However, it is not clear whether this faster absorption translates into better nitrogen utilisation since studies carried out to date have provided conflicting results (Collin-Vidal *et al*; 1994; Endocrinol. Metab., 30, E 907-914). Further, this interest is in the sense of

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providing a source of amino acids to meet the general amino acid needs of the patient and not to specifically provide for the needs of individual organs.

However, in many instances, a person may be suffering from organ specific problems such as depletion of cells, incomplete or improper functioning, underdevelopment or fatigue, in the organs. Providing nutrition to the person in general sense, although beneficial, does not address these organ specific problems.

Various naturally-occurring and synthetic peptides have been reported in the literature as being useful for stimulating the growth of specific tissue. For example, international patent publication WO 92/20707 discloses certain bombesin analogs which may be used to stimulate or antagonise growth of lung tissue. Bombesin itself is a peptide of 14 amino acids which is isolated from the skin of the frog Bombina Bombina. As another example, European patent 0017867 discloses the use of a hydrolysate of a blood plasma extract as an agent for increasing liver growth. Other synthetic peptides for increasing liver growth are disclosed in Pickart *et al*; 1973; Biochem. Biophys. Res. Commun. 54(2), 562-6 (Gly-His-Lys); and Japanese patent application 05-229940 (di- and tri-peptides containing Ala and Gln). However these peptides are not considered dietary protein and it is not practicable to use them as the primary protein source in nutritional formulas.

Summary of the Invention

It has now been discovered that the form in which dietary protein is provided has a selective effect on the protein concentration, RNA concentration, ribosomal efficacy, and rate of protein synthesis in different organs. This offers the advantage of being able to promote the growth or recovery of a particular organ by providing the dietary protein in a form which increases the protein concentration or the rate of protein synthesis, or both, in that organ.

Accordingly, on one aspect, this invention provides a method for promoting the growth or recovery of a selected organ in a mammal, the method comprising administering to the mammal a nutritional formula containing a source of dietary protein in a form for increasing the protein concentration or rate of protein synthesis in the selected organ.

Without wishing to be bound by theory, it is believed that protein hydrolysates having a higher degree of hydrolysis are rapidly digested and

absorbed in the upper small intestine. Therefore protein substrate is available for protein synthesis in the upper small intestine. In this way, the upper small intestine may be targeted. Intact protein and protein hydrolysates having a lower degree of hydrolysis take longer to digest and are more slowly absorbed in the lower small intestine. Therefore protein substrate is available for protein synthesis in the lower small intestine. Also, the lower rate of absorption may result in more protein substrate being available for protein synthesis in the muscles due to the decrease in liver oxidation. In this way, the lower small intestine and muscles may be targeted.

In another aspect, this invention provides the use of a selected form of dietary protein which increases protein concentration or rate of protein synthesis in a selected organ as a protein source in the preparation of a nutritional formula for promoting the growth or recovery of the selected organ in a mammal.

Preferably the source of dietary protein is in the form of a protein hydrolysate, or free amino acids, and combinations thereof. Preferably the protein hydrolysates are hydrolysates of milk protein or a peptide mixture isolated from hydrolysates of milk protein.

In one specific embodiment, this invention provides a method for increasing protein concentration and synthesis in the small intestine, the method comprising administering to a patient an effective amount of a nutritional formula containing a protein hydrolysate having a degree of hydrolysis of at least about 30%. Preferably the protein hydrolysate comprises more than about 30% by weight of di- and tri-peptides. Further, the protein hydrolysate preferably has a non protein nitrogen concentration of at least about 85% of total nitrogen.

Preferably, the method may be used to treat patients suffering from illnesses or damage to the duodenum and jejunum and to promote recovery of the duodenum and jejunum. The method is more preferably used to treat patients suffering from illnesses or damage to the duodenum and to promote recovery of the duodenum.

In another specific embodiment, this invention provides a method for increasing protein concentration and synthesis in the jejunum, the method comprising administering to a patient an effective amount of a nutritional formula containing a protein source in the form of (i) a protein hydrolysate having a degree of hydrolysis of at least about 15%; (ii) free amino acids; or (iii) mixtures thereof. Preferably the protein hydrolysate comprises more than about 20% by weight of di- and tri-peptides. Further, the protein hydrolysate

preferably has a non protein nitrogen concentration of at least about 60% of total nitrogen.

Preferably, the method may be used to treat patients suffering from illnesses or damage to the jejunum to promote recovery of the jejunum.

5 In another aspect, this invention provides a method for the treatment of a patient suffering from illness of or damage to the duodenum, the method comprising (i) administering to a patient an effective amount of a nutritional formula containing a protein hydrolysate having a degree of hydrolysis of at least about 30% to initiate recovery of the duodenum; and (ii) thereafter administering
10 to a patient a nutritional formula containing dietary protein in the form of whole protein or a hydrolysate having a degree of hydrolysis less than about 15%.

In another aspect, this invention provides a method for maintaining muscle protein synthesis and for the prophylaxis or treatment of muscular atrophy, the method comprising administering a patient an effective amount of a nutritional
15 formula containing dietary protein in the form of whole protein, a hydrolysate having a degree of hydrolysis less than about 15%, or free amino acids, or a combination thereof.

The patient may have a compromised gut function and is preferably administered a nutritional formula containing dietary protein in the form of free
20 amino acids.

In another aspect, this invention provides a nutritional formula for increasing protein concentration and synthesis in the intestine, the nutritional formula comprising: a source of dietary protein in the form of a protein hydrolysate having a degree of hydrolysis of at least about 30%, a non protein
25 nitrogen concentration of at least about 85% of total nitrogen, and comprising more than about 30% by weight of di- and tri-peptides; a carbohydrate source; and a lipid source.

Preferably at least about 60% of the components of the protein hydrolysate have a molecular weight less than about 367; more preferably at least about 50%
30 of the components of the protein hydrolysate have a molecular weight of about 127 to about 367.

The nutritional formula is preferably an enterally administrable formula; for example in the form of a powder, a liquid concentrate, or a ready-to-drink beverage.

35 Embodiments of the invention are now described by way of example only.

Detailed Description of Preferred Embodiments of the Invention

In this specification, the term "degree of hydrolysis" (DH) means the percentage of nitrogen in the form of amino nitrogen as compared to total nitrogen. It is a measure of the extent to which the protein has been hydrolysed.

It has been discovered that the form in which dietary protein is consumed influences various organs to different extents. This provides the significant advantage that the growth or recovery of a specific organ may be selectively enhanced through nutrition. All that is required is to specifically enhance the growth or recovery of an organ is administer a nutritional formula which contains the dietary protein in the correct form.

The dietary protein which is used may be any suitable dietary protein; for example animal proteins (such as milk proteins, meat proteins and egg proteins); vegetable proteins (such as soy protein, wheat protein, rice protein, and pea protein); mixtures of free amino acids; or combinations thereof. Milk proteins such as casein and whey protein are particularly preferred.

For targeting nutrition to the small intestine, one suitable form of the dietary protein is protein hydrolysate. Casein and whey protein hydrolysates are preferred. The extent to which the protein is hydrolysed influences the area of the intestine in which the protein is digested and used for protein synthesis.

In particular, hydrolysates having a degree of hydrolysis of about 10% to about 15%, are found to increase relative weight of the liver as compared to free amino acid mixes. Further, these hydrolysates may be used for maintaining muscle protein synthesis. Hydrolysates having a degree of hydrolysis of about 15% to about 25% are found to increase the concentration of protein in the jejunum, the relative weight of the jejunum and the rate of protein synthesis in the jejunum. Highly hydrolysed protein which has a degree of hydrolysis of greater than 25% or which contains more than 25% by weight of di- and tri-peptides, more preferably greater than 30%, is found to increase the rate of protein synthesis in the jejunum and the duodenum; particularly the duodenum.

The protein hydrolysates may be produced using procedures which are well known in the art or may be obtained commercially. For example, nutritional formulas containing hydrolysates having a degree of hydrolysis less than about 15% are commercially available from Nestlé Nutrition Company under the trade mark Peptamen®. Hydrolysates having a degree of hydrolysis above about 15% may be prepared using the procedure described in EP 0322589.

The dietary protein may also be in the form of a mix of free amino acids; preferably such that the mix provides a balanced amino acid profile. Dietary protein in the form of a mix of free amino acids is found to increase the relative weight of the jejunum and the rate of protein synthesis in the jejunum. Further, free amino acids maintain levels of protein synthesis in the muscles while being in an easily digestible form. Hence free amino acids are suitable for maintaining muscle protein synthesis in patients having compromised gut function.

It is also possible to provide the dietary protein in a variety of forms such that several organs are simultaneously provided specific nutrition.

The source of dietary protein preferably provides about 5% to about 30% of the energy of the nutritional formula; for example about 10% to about 20% of the energy. The remaining energy of the nutritional formula may be provided in the form of carbohydrates and fats.

If the nutritional formula includes a fat source, the fat source preferably provides about 5% to about 55% of the energy of the nutritional formula; for example about 20% to about 50% of the energy. The lipids making up the fat source may be any suitable fat or fat mixture. Vegetable fats are particularly suitable; for example soy oil, palm oil, coconut oil, safflower oil, sunflower oil, corn oil, canola oil, lecithins, and the like. Animal fats such as milk fats may also be added if desired. The lipids may also include medium-chain triglycerides; for example up to about 80% by weight of lipids as medium-chain triglycerides. Fractionated coconut oil is a suitable source of medium-chain triglycerides.

If the nutritional formula includes a carbohydrate source, the carbohydrate source preferably provides about 40% to about 80% of the energy of the nutritional formula. Any suitable carbohydrates may be used, for example sucrose, lactose, glucose, fructose, corn syrup solids, and maltodextrins, and mixtures thereof.

Dietary fibre may also be added if desired. Numerous types of dietary fibre are available. Suitable sources of dietary fibre, among others, include soy, pea, oat, pectin, guar gum, and gum arabic. If used, the dietary fibre preferably comprises up to about 5% of the weight of the nutritional formula.

Suitable vitamins and minerals may be included in the nutritional formula in the usual manner to meet the appropriate guidelines.

One or more food grade emulsifiers may be incorporated into the nutritional formula if desired; for example diacetyl tartaric acid esters of mono-diglycerides, lecithin and mono- and di-glycerides. Similarly suitable salts and stabilisers may be included.

The nutritional formula may be prepared in any suitable manner. For example, the nutritional formula may be prepared by blending together the source of dietary protein, the carbohydrate source, and the fat source in appropriate proportions. If used, the emulsifiers may be included in the blend. The vitamins and minerals may be added at this point but are usually added later to avoid thermal degradation. Any lipophilic vitamins, emulsifiers and the like may be dissolved into the fat source prior to blending. Water, preferably water which has been subjected to reverse osmosis, may then be mixed in to form a liquid mixture. The temperature of the water is conveniently about 50°C to about 80°C to aid dispersal of the ingredients. Commercially available liquefiers may be used to form the liquid mixture. The liquid mixture is then homogenised; for example in two stages.

The liquid mixture may then be thermally treated to reduce bacterial loads. For example, the liquid mixture may be rapidly heated to a temperature in the range of about 80°C to about 150°C for about 5 seconds to about 5 minutes. This may be carried out by steam injection, autoclave or by heat exchanger; for example a plate heat exchanger.

The liquid mixture may then be cooled to about 60°C to about 85°C; for example by flash cooling. The liquid mixture may then be again homogenised; for example in two stages at about 7 MPa to about 40 MPa in the first stage and about 2 MPa to about 14 MPa in the second stage. The homogenised mixture may then be further cooled to add any heat sensitive components; such as vitamins and minerals. The pH and solids content of the homogenised mixture is conveniently standardised at this point.

If it is desired to produce a powdered nutritional formula, the homogenised mixture is transferred to a suitable drying apparatus such as a spray drier or freeze drier and converted to powder. The powder should have a moisture content of less than about 5% by weight.

If it is desired to produce a liquid formula, the homogenised mixture is preferably aseptically filled into suitable containers. Aseptic filling of the containers may be carried out by pre-heating the homogenised mixture (for example to about 75 to 85°C) and then injecting steam into the homogenised mixture to raise the temperature to about 140 to 160°C; for example at about 150°C. The homogenised mixture may then be cooled, for example by flash cooling, to a temperature of about 75 to 85°C. The homogenised mixture may then be homogenised, further cooled to about room temperature and filled into

containers. Suitable apparatus for carrying out aseptic filling of this nature is commercially available. The liquid formula may be in the form of a ready to feed formula having a solids content of about 10 to about 14% by weight or may be in the form of a concentrate; usually of solids content of about 20 to about 26% by weight. Flavours may be added to the liquid formulas so that the formulas are provided in the form of convenient, flavoursome, ready-to-drink beverages.

The nutritional formula may be used in the prophylaxis or treatment of a variety of conditions or diseased states. For example, many premature babies suffer from undeveloped intestines. A nutritional formula containing dietary protein in the form of hydrolysed protein or free amino acids may be fed to the babies to promote an increase in protein concentration and protein synthesis in intestinal tissue and hence increase development of the intestine. As a further example, many elderly people suffer from atrophy of the intestine. Hence these nutritional formulas may be fed to the elderly for the prophylaxis or treatment of intestinal atrophy.

Further, nutritional formulas containing dietary protein in the form of hydrolysed protein or free amino acids may be fed to patients who suffer from illnesses or damage to the intestine to promote recovery of the intestine; for example, inflammatory conditions of the gastro-intestinal tract (such as Crohn's disease and sepsis), gut epithelial damage, consequences of severe diarrhoea, post antibiotic colitis, gut surgery, jejunum resection, and atrophy after parenteral feeding.

The nutritional formulas may also be used to assist in the recovery of muscles after exercise or in the prophylaxis or treatment of muscular atrophy in the elderly. In this case, the source of dietary protein may be in the form of free amino acids, intact protein, or a hydrolysate having a degree of hydrolysis of less than about 15%.

The amount of the nutritional formula to be administered will vary depending upon the age of the patient, the condition or disease, and the severity of the condition or disease. However the amount may be readily set by a medical practitioner. In general, an amount of the nutritional formula sufficient to provide a daily dosage of dietary protein of about 3 g to about 300 g. A daily dosage of dietary protein of about 50 g to about 150 g is preferred for clinical applications.

The nutritional formula may be taken in multiple doses, for example 2 to 10 times a day, to make up the selected daily dosage or may taken in a single dose. If in a single dose form, the nutritional formula is conveniently taken in replacement of a meal. In the case of multiple doses, the nutritional formula is conveniently in the form of a convenience food; for example a ready-to-drink beverage. The nutritional formula may also be administered continuously by means of nasogastric tubes or enteral tubes such as jejunum tubes.

Example 1

a) Whole proteins

An amount of 5 kg of whey protein (obtained from Meggle GmbH under the trade name Globulal 80) is dispersed in demineralised water at 55°C to obtain protein concentration (N*6.38) of 10% by weight. The pH of the dispersion is adjusted by the addition of 190 g of calcium hydroxide and the dispersion is cooled to room temperature. The proteins are then dried by lyophilisation and packaged into metal cans.

The whole proteins have a degree of hydrolysis of about 4.41% and a non protein nitrogen concentration of about 1.1% on the basis of total nitrogen.

b) Hydrolysate 1

An amount of 6.25 kg of whey protein (obtained from Meggle GmbH under the trade name Globulal 80) is dispersed in 50 litres of demineralised water at 55°C. The pH of the dispersion is adjusted to 8.2 by the addition of 1.8 litres of 2M $\text{Ca}(\text{OH})_2$. The proteins are then hydrolysed using 30 g of trypsin (Salt free pancreatic trypsin which has an activity of 6.8 AU/g and a chymotrypsin content of less than 5% and which is obtainable from Novo Nordisk Ferment AG, Dittigen, Switzerland). The hydrolysis reaction is continued for 4 hours at 55°C. During the reaction, the pH is regulated to 7.4 by the addition of 1.6N NaOH and 0.4N KOH. The enzymes are then inactivated by heating the reaction mixture to 80°C and holding the mixture at this temperature for about 5 minutes. The mixture is then cooled to 16°C. The hydrolysed proteins are then dried by lyophilisation and packaged into metal cans. A sample of the hydrolysed

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proteins is then subjected to HPLC gel filtration to determine the molecular weight distribution.

Molecular weight	Percentage	Cumulative percentage
33730	0	0
25820	2.3	2.3
19860	4.3	6.6
15210	5.5	12.1
11670	6.9	19.0
8933	3.9	22.9
6855	3.6	26.5
5248	3.1	29.6
4027	3.7	33.3
3083	4.5	37.8
2366	9.4	47.2
1811	6.9	54.1
1387	6.9	61.0
1064	3.9	64.9
815	9.2	74.2
625	6.7	80.9
479	1.6	82.5
367	1.5	84.0
281	3.6	87.6
215	8.4	96.0
165	3.4	99.4
127	0.4	99.8
97	0.1	99.9
74	0.1	100.0

- 5 The hydrolysate has a degree of hydrolysis of about 14% and a non protein nitrogen concentration of about 54.5% on the basis of total nitrogen.

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c) Hydrolysate 2

An amount of 6.25 kg of whey protein (obtained from Meggle GmbH under the trade name Globulal 80) is dispersed in 50 litres of demineralised water at 55°C. The pH of the dispersion is adjusted to 7.5 by the addition of 1.6 litres of 1M Ca(OH)₂ and 162 ml of a solution of 1.6M NaOH and 0.4M KOH. The proteins are then hydrolysed using 50 g of trypsin (obtainable from Novo Nordisk Ferment AG). The hydrolysis reaction is continued for 4 hours at 55°C. During the reaction, the pH is regulated to 7.4 by the addition of 1.6N NaOH and 0.4N KOH. The enzymes are then inactivated and non-hydrolysed protein is denatured, by heating the reaction mixture to 90°C and holding the mixture at this temperature for about 5 minutes.

The mixture is then cooled to 56°C and hydrolysed again for 1 hour using 50g of trypsin at 55°C. During the reaction, the pH is regulated to 7.4 by the addition of 1.6N NaOH and 0.4N KOH. The enzymes are then inactivated by heating the reaction mixture to 80°C and holding the mixture at this temperature for about 5 minutes. The mixture is then cooled to 18°C. The hydrolysed proteins are then dried by lyophilisation and packaged into metal cans.

A sample of the hydrolysed proteins is then subjected to HPLC gel filtration to determine the molecular weight distribution.

Molecular weight	Percentage	Cumulative percentage
33730	1.0	1.0
25820	2.7	3.7
19860	3.0	6.7
15240	2.8	9.5
11670	3.2	12.7
8954	2.1	14.8
6855	2.2	16.9
5260	2.3	19.2
4027	3.0	22.2
3090	3.8	26.0
2366	9.7	35.7
1811	7.0	42.7

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1390	7.5	50.2
1064	4.2	54.4
817	10.4	64.8
625	8.7	73.5
480	2.1	75.6
367	1.7	77.3
282	5.2	82.5
216	11.1	93.6
165	4.9	98.5
127	0.8	99.3
97	0.2	99.5
74	0.3	99.8
57	0.2	100.0

The hydrolysate has a degree of hydrolysis of about 17.3% and a non protein nitrogen concentration of about 65.9% on the basis of total nitrogen.

5 d) Hydrolysate 3

An amount of 6.25 kg of whey protein (obtained from Meggle GmbH under the trade name Globulal 80) is dispersed in 50 litres of demineralised water at 55°C. The pH of the dispersion is adjusted to 7.5 by the addition of 1.6 litres of 10 1M Ca(OH)₂ and 162 ml of a solution of 1.6M NaOH and 0.4M KOH. The proteins are then hydrolysed using 250 g of Alcalase 2.4L (EC 940459 - obtainable from Novo Nordisk Ferment AG). The hydrolysis reaction is continued for 4 hours at 55°C. For the first hour of the reaction, the pH is regulated to 7.6 by the addition of 1.6N NaOH and 0.4N KOH.

15 An amount of 250g of Neutrase 0.5L (obtainable from Novo Nordisk Ferment AG) is added and the proteins are further hydrolysed for 4 hours at 50°C. The enzymes are then inactivated by heating the reaction mixture to 90°C and holding the mixture at this temperature for about 5 minutes. The reaction mixture is then cooled to 55°C.

20 The pH of the reaction mixture is adjusted to 7.33 by the addition of 1.6N NaOH and 0.4N KOH and the reaction mixture hydrolysed again for 4 hours using 100g of pancreatin at 55°C. During the reaction, the pH is regulated to 7.5

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by the addition of 1M NaOH. The enzymes are then inactivated by heating the reaction mixture to 90°C and holding the mixture at this temperature for about 5 minutes. The mixture is then cooled to 4°C. The hydrolysed proteins are then dried by lyophilisation and packaged into metal cans.

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A sample of the hydrolysed proteins is then subjected to HPLC gel filtration to determine the molecular weight distribution.

Molecular weight (Dalton)	Percentage	Cumulative percentage
5248	0.1	0.1
3083	0.1	0.2
2366	0.9	1.1
1811	1.2	2.3
1387	3.0	5.3
1064	1.4	6.7
815	6.7	13.4
625	14.2	27.6
479	5.3	32.9
367	1.6	34.5
281	10.3	44.8
215	22.2	67.0
165	14.5	82.4
127	6.8	89.2
97	10.5	99.7
74	0.2	99.9
57	0.1	100.00

10

The hydrolysate has a degree of hydrolysis of about 35% and a non protein nitrogen concentration of about 92.6% on the basis of total nitrogen. The hydrolysate is rich in small peptides, especially di- and tri-peptides which have a molecular weight of about 127 to 367.

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Example 2

a) Experimental Feeds

- 5 Five different feeds are used in the trials. The overall composition of these feeds is given in table 1.

Table 1: Feed composition

	Feed 1 (g/100g)	Feed 2 (g/100g)	Feed 3 (g/100g)	Feed 4 (g/100g)	Feed 5 (g/100g)
Protein source	14.62	14.77	14.78	14.93	15.51
Carbohydrate	64.7	64.4	64.1	63.4	65.4
Lipids	8.21	8.27	8.26	8.37	7.90
Minerals	6.18	5.72	5.69	5.07	6.38
Ash	2.98	3.00	3.00	3.04	2.87

10

The values are quoted as g of component per 100g of dry matter. The energy provided by all feeds is from 405.9 to 412.5 kCal/100 g. The protein sources used in each feed are different and are as given in table 2.

15

Table 2 - Protein Source for feeds

Feed	Protein Source
1	whole protein from example 1
2	Hydrolysate 1 from example 1
3	Hydrolysate 2 from example 1
4	Hydrolysate 3 from example 1
5	Free amino acids

The amino acid composition of each protein source is given in table 3.

Table 3 - Amino Acid Composition of Protein Source

Amino Acid	Feed 1 (g/100g)	Feed 2 (g/100g)	Feed 3 (g/100g)	Feed 4 (g/100g)	Feed 5 (g/100g)
Asp	1.00	1.01	1.01	1.02	1.06
Asn	0.66	0.67	0.67	0.68	0.71
Thr	0.77	0.78	0.78	0.79	0.82
Ser	0.72	0.73	0.73	0.73	0.76
Glu	1.60	1.62	1.62	1.64	1.70
Gln	0.86	0.87	0.87	0.88	0.91
Pro	0.60	0.61	0.61	0.62	0.64
Gly	0.29	0.30	0.30	0.30	0.31
Ala	0.68	0.69	0.69	0.70	0.72
Val	0.71	0.72	0.72	0.72	0.75
Cys ¹	0.43	0.43	0.43	0.43	0.45
Met	0.29	0.30	0.30	0.30	0.31
Ile	0.74	0.74	0.74	0.75	0.78
Leu	1.80	1.82	1.82	1.84	1.91
Tyr	0.51	0.51	0.51	0.52	0.54
Phe	0.53	0.54	0.54	0.54	0.56
Lys	1.43	1.44	1.44	1.46	1.51
His	0.27	0.28	0.28	0.28	0.29
Arg	0.42	0.42	0.42	0.43	0.44
Trp	0.31	0.31	0.31	0.32	0.33
Total amino acid	14.62	14.77	14.78	14.93	15.51

(1) in the form of cystine for Feed 5.

5

b) Experimental Protocol and weight determination

A group of 40 male, Sprague Dawley rats, supplied by IFFA CREDO of l'Arbresle, France are used. The rats have an average weight of $82.3 \text{ g} \pm 1.0$.

10 The rats are lodged in separate cages which are illuminated from 7.30 am to 7

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pm. For a few days prior to the trials, the rats are allowed to feed *ad libitum* on a standard, pelleted food. The rats have free access to water.

Two days prior to the trials, the rats are separated into 5 groups of 8 rats. Each group of rats is fed, *ad libitum*, one of feeds 1 to 5 for an adaptation period of 2 days. Starting with the first day of the trial, each day at 3 am, 7 am, 11 am, 3 pm, 7 pm and 11 pm, the rats of each group are then fed a precisely metered amount of the feed which they had received during the adaptation period. The feeding regime is continued for a period of 15 days. Each rat is weighed each day.

On the 16th day, a radioactive mixture is prepared by adding 1.7 g of a solution containing ^{14}C Phe (170 μCi per 2.9 μmol) to 9.3 g of a solution containing 30.2 g/l Phe (non radioactive). Half of the rats in each group are anaesthetised and slaughtered at 1.30 pm; the other half are anaesthetised and slaughtered at 3 pm. About 20 minutes prior to being slaughtered, the radioactive mixture is injected intravenously into each rat at 2 ml/200g of body weight.

After slaughter, the blood of each rat is removed and immediately centrifuged at 4°C. The plasma is then separated into fractions and the fractions stored at -20°C. The liver of each rat is rapidly removed, washed with a cold solution containing 9% NaCl, dried, weighed, and frozen in liquid nitrogen. The intestine of each rat is removed, emptied and rinsed in a 5% solution of trichloroacetic acid and separated into three fractions; the whole duodenum, about 13 cm of the jejunum and the remainder of the intestine. Each fraction is dried, weighed, and frozen in liquid nitrogen. The fractions are ground in the liquid nitrogen prior to use. The stomach is removed and weighed before and after emptying. The muscles (gastrocnemius, soleus, and extensor digitorum longus) are removed, de-fatted, washed, dried, weighed and frozen in liquid nitrogen.

The mean weight of the rats at slaughter and the mean weight of the body parts are as follows:

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		Feed 1	Feed 2	Feed 3	Feed 4	Feed 5
Body weight/g		173.9	159.4	164.2	154.1	157.7
Stomach	Absolute g	1.19	1.11	1.32	1.25	1.32
	Relative %	0.68	0.70	0.78	0.80	0.84
Intestine	Absolute g	5.95	5.51	5.66	6.02	5.65
	Relative %	3.42	3.46	3.44	3.90	3.59
Duodenum	Absolute g	0.40	0.34	0.34	0.36	0.38
	Relative %	0.23	0.21	0.21	0.24	0.24
Jejunum	Absolute g	1.13	1.15	1.24	1.21	1.24
	Relative %	0.65	0.73	0.76	0.78	0.79
Liver	Absolute g	8.59	8.40	7.87	7.37	7.29
	Relative %	4.93	5.24	4.77	4.80	4.62
Gastrocnemius						
	Absolute g	0.96	0.96	1.04	0.95	0.87
	Relative %	0.55	0.60	0.64	0.62	0.55
Soleus	Absolute g	0.069	0.066	0.068	0.064	0.060
	Relative %	0.040	0.041	0.042	0.042	0.038
Extensor	Absolute g	0.075	0.075	0.072	0.070	0.067
	Relative %	0.043	0.047	0.044	0.046	0.042

The absolute value is the weight of the organ in grams. The relative value is a percentage based on the weight of the organ per 100g of body weight.

- 5 The rats fed with Feeds 1 and 3 have a noticeably higher body weight than those feed with the other feeds. The relative weight of the whole intestine of the rats is significantly ($P<0.05$) higher with Feed 4 than the other feeds. Also, the relative weight of the jejunum of the rats is significantly ($P<0.05$) higher with Feeds 3, 4 and 5 than the other feeds. The weight of the liver of the rats fed with
- 10 Feed 5 is significantly lower ($P<0.05$) than with the other feeds.

c) Determination of protein and RNA content

- 15 Tissue samples of 1 g of muscle, 1 g of liver, and 0.5 to 1 g intestine are extracted with a cold solution of 10% trichloroacetic acid to dissolve free amino acids and precipitate protein. The extract is centrifuged at 4°C and 10000 g for

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15 minutes. The residue is extracted with a cold solution of 10% trichloroacetic acid and centrifuged at 4°C and 10000 g for 15 minutes. The extraction and centrifuging procedure is repeated again. The residue is collected and the three filtrates are combined.

- 5 A column containing 3 ml of Amberlite resin (AG 50 X8, 100-200 mesh) in H⁺ form is neutralised to fix free amino acids. The combined filtrate is added to the column and the column washed three times with distilled water. Free amino acids are then eluted using 4N ammonia. The eluant is collected and the ammonia removed by evaporation under vacuum at 40°C. The dry residue is rehydrated to provide a first amino acid solution having a Phe concentration in the region of about 1 µmol/ml. The pH of the first amino acid solution is adjusted to 7.

- 15 The residue from the centrifuge is extracted with cold 0.2 M perchloric acid and the extract is centrifuged at 4°C and 10000 g for 15 minutes. The residue is again extracted with perchloric acid and centrifuged in the same manner. The residue is then rehydrated in a solution of water and 3N NaOH. An aliquot of the rehydrated residue is collected for quantification of the protein using the bicinchoninic acid method described in Smith *et al*; 1985; Anal. Biochem., 150, 76-85.

- 20 An amount of 3N perchloric acid is added to the remaining rehydrated residue to precipitate protein. The solution is centrifuged at 4°C at 15000 g for 10 minutes. The RNA concentration in the filtrate is then determined by spectrophotometry ($\lambda_1=232$ and $\lambda_2=260$) using the technique of Baillie A.G.S and Garlick, P.J.; 1991; Am. J. Physiol., 260, E891-E896.

- 25 The residue from the centrifuge is hydrolysed using 5.5N HCl at 150°C over 24 hours to liberate Phe. The HCl is evaporated under vacuum and the residue is rehydrated with water to provide a second amino acid solution having a Phe concentration of about 1 µmol/ml. The pH of the second amino acid solution is adjusted to 7.

30

d) Determination of Phe radioactivity and rate of Protein synthesis

- 35 A Kontron HPLC and Kontron U.V. spectrophotometer ($\lambda = 330\text{nm}$) are used to separate and detect amino acids. Two solvents are used for the mobile phase:

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- solvent A comprising 69g of $\text{NaH}_2\text{PO}_4 \cdot \text{H}_2\text{O}$, 32 ml of H_3PO_4 , 40g of NaOH particles, a 30% NaOH solution to adjust the pH to 7, and ultra pure water to adjust the volume to 5 litres.

5 - solvent B comprising 200 ml of solvent A diluted with 300 ml of ultra pure water, 350 ml of methanol, and 150 ml of acetonitrile.

i) Radioactivity of free Phe

10 The HPLC column is fitted with a Spherisorb OD2 column (5 μm , 250 x 4.6mm) and is operated at 25°C. An orthophthaldialdehyde solution is added to the first amino acid solution in a mass ratio of 2:1 and the mixture is allowed to react. An amount of 40 μl of the mixture is added to the column. The column is eluted with solvents A and B at various times; the total amount being added at each time being 1 ml/min. The ratio of solvent A to solvent B at each time is
15 100:0 at time 0, 40:60 after 7 minutes, 27:73 after 22 minutes, 0:100 after 24 minutes, 0:100 after 26 minutes, and 100:0 after 28 minutes. Elution is continued for 33 minutes. The eluant is collected in fractions of 0.5 ml. Standard solutions containing a mixture of 20 amino acids, including ^{14}C -Phe, are also run through the column to identify and standardise the amino acid peaks.

20 The fractions of the first amino acid solution containing the Phe peak are collected and the volume adjusted to 5 ml with ultra pure water. Ten ml of a scintillant (Quick Safe A) is added to the solution. The radioactivity is then determined using a Kontron scintillation detector.

25 ii) Radioactivity of bound Phe

30 The HPLC column is fitted with a Spherisorb OD2 column (5 μm , 250 x 10mm) and is operated at 25°C. An orthophthaldialdehyde solution is added to the second amino acid solution in a mass ratio of 1:1 and the mixture is allowed to react. An amount of 800 μl of the mixture is added to the column. The column is eluted with solvents A and B at various times; the total amount being added at each time being 4.25 ml/min. The ratio of solvent A to solvent B at each time is
35 100:0 at time 0, 50:50 after 7 minutes, 34:66 after 27 minutes, 0:100 after 29 minutes, 0:100 after 32 minutes, and 100:0 after 34 minutes. Elution is continued for 39 minutes. The eluant is collected in fractions of 0.5 ml.

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Standard solutions containing a mixture of 20 amino acids, including ^{14}C -Phe, are also run through the column to identify and standardise the amino acid peaks.

- 5 The fractions of the second amino acid solution containing the Phe peak are collected and the volume adjusted to 5 ml with ultra pure water. Ten ml of a scintillant (OPTI Phase "Hisafe") is added to the solution. The radioactivity is then determined using a Kontron scintillation detector.

iii) Calculation of the rate of protein synthesis

- 10 A measure of the rate of protein synthesis may be obtained using the formula (McNurlan *et al*; 1979; Biochem. J. 178, 373-379):

$$\text{Rate of synthesis (\% per day)} = S_B / (S_A \cdot t)$$

- 15 Where S_B is the specific radioactivity value (dpm/ μmol) of the bound Phe; S_A is the mean specific radioactivity value (dpm/ μmol) of the free Phe; and t is the time in days.

- 20 e) Protein and RNA content and rate of protein synthesis.

The protein and RNA content of the different organs and the rate of protein synthesis in the different organs are as follows for each feed:

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<u>Liver</u>	Feed 1	Feed 2	Feed 3	Feed 4	Feed 5
Protein Concentration					
mg/g tissue	139.9	160.9	156.4	165.2	147.2
g/100g rat	0.69	0.84	0.75	0.79	0.68
Total protein, g	1.20	1.34	1.24	1.23	1.07
RNA Concentration					
mg/g tissue	7.89	8.10	8.00	8.20	8.17
mg/100g rat	38.8	42.4	38.1	38.9	37.7
Total RNA, mg	67.47	67.93	62.73	60.43	59.45
Protein Synthesis Capacity (mg RNA/g protein)	56.46	51.86	51.76	50.62	56.62
Protein Synthesis Rate (% per day)	82.29	86.94	85.55	89.68	91.53
Daily Protein Synthesis (g protein/day)	1.01	1.18	1.08	1.16	1.01
Ribosomal Efficacy (mg protein/day.mgRNA)	14.69	17.53	17.00	18.50	16.85

No significant difference between the effects that the feeds have on the liver are noticed. Feed 2 however appears to provide a larger relative concentration of protein and RNA in the liver.

5

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Intestine (Duodenum)	Feed 1	Feed 2	Feed 3	Feed 4	Feed 5
Protein Concentration					
mg/g tissue	178.6	180.9	165.0	191.4	162.0
g/100g rat	41.9	39.1	34.2	46.4	41.1
Total protein, g	72.0	61.8	56.0	71.3	65.0
RNA Concentration					
mg/g tissue	8.32	7.97	6.55	8.43	7.59
mg/100g rat	1.98	1.64	1.36	2.05	1.92
Total RNA, mg	3.39	2.59	2.21	3.12	3.03
Protein Synthesis Capacity (mg RNA/g protein)	46.23	44.46	39.69	44.23	48.03
Protein Synthesis Rate (% per day)	97.9	105.7	100.3	135.9	108.0
Daily Protein Synthesis (g protein/day)	72.99	68.52	57.01	97.89	74.44
Ribosomal Efficacy (mg protein/day.mgRNA)	21.2	25.7	25.3	30.8	23.0

5 Feed 4 has a significant effect ($P < 0.05$) on the protein synthesis rate in the duodenum as compared to the other feeds. Protein and RNA concentrations are raised but not significantly. Further the daily protein synthesis and the ribosomal efficacy are all raised.

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<u>Intestine (Jejunum)</u>	Feed 1	Feed 2	Feed 3	Feed 4	Feed 5
Protein Concentration					
mg/g tissue	109.9	109.5	112.2	106.5	110.0
g/100g rat	71.7	79.6	84.1	83.8	87.1
Total protein, g	124.8	126.3	138.7	130.2	137.2
RNA Concentration					
mg/g tissue	7.56	7.39	7.55	7.36	7.87
mg/100g rat	4.90	5.38	5.71	5.82	6.23
Total RNA, mg	8.49	8.50	9.34	8.98	9.79
Protein Synthesis Capacity (mg RNA/g protein)	70.35	67.48	68.31	69.58	71.70
Protein Synthesis Rate (% per day)	89.9	88.4	103.8	111.8	109.5
Daily Protein Synthesis (g protein/day)	111.6	111.8	141.4	144.9	147.7
Ribosomal Efficacy (mg protein/day.mgRNA)	13.02	13.25	15.15	16.14	15.24

The rate of protein synthesis, the daily protein synthesis, and the ribosomal efficacy are significantly ($P < 0.05$) raised with feeds 3 to 5 as compared to feeds 1 and 2.

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Muscle (Gastrocnemius)	Feed 1	Feed 2	Feed 3	Feed 4	Feed 5
Protein Concentration					
mg/g tissue	189.0	183.2	189.8	180.7	182.4
g/100g rat	109.3	109.7	113.2	108.4	99.9
Total protein, g	188.2	174.7	185.1	166.4	157.4
RNA Concentration					
mg/g tissue	2.14	2.18	1.97	1.91	1.95
mg/100g rat	1.24	1.31	1.18	1.15	1.07
Total RNA, mg	2.14	2.10	1.92	1.77	1.69
Protein Synthesis Capacity (mg RNA/g protein)	11.31	11.94	10.36	10.59	10.84
Protein Synthesis Rate (% per day)	11.00	9.87	9.53	8.83	11.26
Daily Protein Synthesis (g protein/day)	20.75	17.33	18.01	15.38	17.58
Ribosomal Efficacy (mg protein/day.mgRNA)	9.69	8.57	9.17	8.24	10.33

No significant differences are determined. However the daily protein synthesis is reduced with feed 4 when compared to feeds 1 and 5.

Claims

- Article
34
1. Use of a selected form of dietary protein which increases protein concentration or rate of protein synthesis in a selected organ as a protein source
5 in the preparation of a nutritional formula for promoting the growth or recovery of the specific organ in a mammal.
 2. Use according to claim 1 in which the dietary protein is a protein hydrolysate having a degree of hydrolysis of at least about 30% for the
10 preparation of a nutritional formula for increasing protein concentration and synthesis in the small intestine.
 3. Use according to claim 2 in which the dietary protein is used in the preparation of a nutritional formula for increasing protein concentration and
15 synthesis in the duodenum.
 4. Use according to claim 2 or claim 3 in which the protein hydrolysate comprises more than about 30% by weight of di- and tri-peptides and has a non protein nitrogen concentration of at least about 85% of total nitrogen.
20
 5. Use according to claim 1 in which the dietary protein is (i) a protein hydrolysate having a degree of hydrolysis of at least about 15%; (ii) free amino acids; or (iii) mixtures thereof, for the preparation of a nutritional formula for increasing protein concentration and synthesis in the jejunum.
25
 6. Use according to claim 5 in which the dietary protein is a protein hydrolysate which comprises more than about 20% by weight of di- and tri-peptides and which has a non protein nitrogen concentration of at least about 60% of total nitrogen.
30
 7. Use according to claim 1 in which the dietary protein is in the form of free amino acids for the preparation of a nutritional formula for maintaining muscle protein synthesis and for the prophylaxis or treatment of muscular atrophy.

8. Use according to claim 7 in which the dietary protein is used in the preparation of a nutritional formula for mammals having compromised gut function.
- 5 9. Use according to claim 1 in which the dietary protein is a protein hydrolysate for the preparation of a nutritional formula for increasing protein concentration and synthesis in underdeveloped intestines of premature babies.
- 10 10. Use according to claim 9 in which the protein hydrolysate comprises more than about 30% by weight of di- and tri-peptides and has a non protein nitrogen concentration of at least about 85% of total nitrogen.

NO 5986/05

COMBINED DECLARATION FOR PATENT APPLICATION AND POWER OF ATTORNEY

(Includes Reference to PCT International Applications)

ATTORNEY'S
DOCKET NUMBER
P00,0164

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name,
I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint
inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on
the invention entitled:

the specification of which (check only one item below):

- ☐ is attached hereto.
- ☐ was filed as United States application
Serial No. _____
on _____,
and was amended
on _____ (if applicable).
- ☒ was filed as PCT international application
Number PCT/EP98/05843
on September 14, 1998,
and was amended under PCT Article 19
on September 18, 1999 (if applicable).

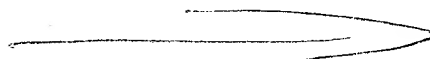
I hereby state that I have reviewed and understand the contents of the above-identified specification, including the
claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance
with Title 37, Code of Federal Regulations, §1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, §119 of any foreign application(s) for
patent or inventor's certificate or of any PCT international application(s) designating at least one country other than
the United States of America listed below and have also identified below any foreign application(s) for patent or
inventor's certificate or any PCT international application(s) designating at least one country other than the United
States of America filed by me on the same subject matter having a filing date before that of the application(s) of
which priority is claimed:

PRIOR FOREIGN/PCT APPLICATION(S) AND ANY PRIORITY CLAIMS UNDER 35 U.S.C. 119:

COUNTRY (if PCT indicate "PCT")	APPLICATION NUMBER	DATE OF FILING (day, month, year)	PRIORITY CLAIMED UNDER 35 USC 119
Europe	97202853.4	September 16, 1997	<input type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO
			<input type="checkbox"/> YES <input type="checkbox"/> NO



**Combined Declaration For Patent Application and Power of Attorney
(Continued)** (Includes Reference to PCT International Applications)ATTORNEY'S DOCKET NO.
P00,0164

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) or PCT international application(s) designating the United States of America that is/are listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in that/those prior application(s) in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, §1.56(a) which occurred between the filing date of the prior application(s) and the national or PCT international filing date of this application:

PRIOR U.S. APPLICATIONS OR PCT INTERNATIONAL APPLICATIONS DESIGNATING THE U.S. FOR BENEFIT UNDER 35 U.S.C. 120:

U.S. APPLICATIONS			STATUS (Check one)		
U.S. APPLICATION NUMBER	U.S. FILING DATE		PATENTED	PENDING	ABANDONED
PCT APPLICATIONS DESIGNATING THE U.S.					
PCT APPLICATION NO	PCT FILING DATE	U.S. SERIAL NUMBERS ASSIGNED (if any)			

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	RESIDENCE & CITIZENSHIP	CITY <u>Corent</u>	STATE OR FOREIGN COUNTRY <u>France FRX</u>	COUNTRY OF CITIZENSHIP <u>France</u>
	POST OFFICE ADDRESS	POST OFFICE ADDRESS <u>7, rue Haute</u>	CITY <u>Corent</u>	STATE & ZIP CODE/COUNTRY <u>France F-63730</u>

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

SIGNATURE OF INVENTOR 201

SIGNATURE OF INVENTOR 202

SIGNATURE OF INVENTOR 203

DATE

March 7th 2000

DATE

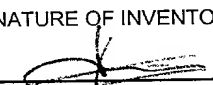
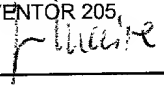
March 13th

DATE

March 1st 2000

Combined Declaration For Patent Application and Power of Attorney (Continued)				ATTORNEY'S DOCKET NO.
400 2 0 4	FULL NAME OF INVENTOR	FAMILY NAME <u>Grizard</u>	FIRST GIVEN NAME <u>Jean</u>	SECOND GIVEN NAME
	RESIDENCE & CITIZENSHIP	CITY <u>Chamalières</u>	STATE OR FOREIGN COUNTRY <u>France</u> <u>FR</u>	COUNTRY OF CITIZENSHIP <u>France</u>
	POST OFFICE ADDRESS	POST OFFICE ADDRESS <u>46, rue des Montagnards</u>	CITY <u>Chamalières</u>	STATE & ZIP CODE/COUNTRY <u>France F-63400</u>
500 2 0 5	FULL NAME OF INVENTOR	FAMILY NAME <u>Maire</u>	FIRST GIVEN NAME <u>Jean-Claude</u>	SECOND GIVEN NAME
	RESIDENCE & CITIZENSHIP	CITY <u>Belmont sur Lausanne</u>	STATE OR FOREIGN COUNTRY <u>Switzerland</u> <u>CH</u>	COUNTRY OF CITIZENSHIP <u>Switzerland</u>
	POST OFFICE ADDRESS	POST OFFICE ADDRESS <u>1, chemin de Rueyres</u>	CITY <u>Belmont sur Lausanne</u>	STATE & ZIP CODE/COUNTRY <u>Switzerland CH-1092</u>
2 0 6	FULL NAME OF INVENTOR	FAMILY NAME	FIRST GIVEN NAME	SECOND GIVEN NAME
	RESIDENCE & CITIZENSHIP	CITY	STATE OR FOREIGN COUNTRY	COUNTRY OF CITIZENSHIP
	POST OFFICE ADDRESS	POST OFFICE ADDRESS	CITY	STATE & ZIP CODE/COUNTRY
2 0 7	FULL NAME OF INVENTOR	FAMILY NAME	FIRST GIVEN NAME	SECOND GIVEN NAME
	RESIDENCE & CITIZENSHIP	CITY	STATE OR FOREIGN COUNTRY	COUNTRY OF CITIZENSHIP
	POST OFFICE ADDRESS	POST OFFICE ADDRESS	CITY	STATE & ZIP CODE/COUNTRY
2 0 8	FULL NAME OF INVENTOR	FAMILY NAME	FIRST GIVEN NAME	SECOND GIVEN NAME
	RESIDENCE & CITIZENSHIP	CITY	STATE OR FOREIGN COUNTRY	COUNTRY OF CITIZENSHIP
	POST OFFICE ADDRESS	POST OFFICE ADDRESS	CITY	STATE & ZIP CODE/COUNTRY
2 0 9	FULL NAME OF INVENTOR	FAMILY NAME	FIRST GIVEN NAME	SECOND GIVEN NAME
	RESIDENCE & CITIZENSHIP	CITY	STATE OR FOREIGN COUNTRY	COUNTRY OF CITIZENSHIP
	POST OFFICE ADDRESS	POST OFFICE ADDRESS	CITY	STATE & ZIP CODE/COUNTRY

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

SIGNATURE OF INVENTOR 204 	SIGNATURE OF INVENTOR 205 	SIGNATURE OF INVENTOR 206
DATE <u>Feb. 21th 2000</u>	DATE <u>March 13, 2000</u>	DATE
SIGNATURE OF INVENTOR 207	SIGNATURE OF INVENTOR 208	SIGNATURE OF INVENTOR 209
DATE	DATE	DATE